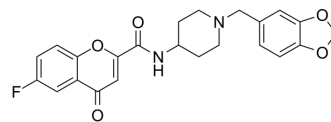


## MCHR1 antagonist 2

<b>Cat. No.:</b>	HY-100321		
<b>CAS No.:</b>	863115-70-2		
<b>Molecular Formula:</b>	C <sub>23</sub> H <sub>21</sub> FN <sub>2</sub> O <sub>5</sub>		
<b>Molecular Weight:</b>	424.42		
<b>Target:</b>	MCHR1 (GPR24); Potassium Channel		
<b>Pathway:</b>	GPCR/G Protein; Neuronal Signaling; Membrane Transporter/Ion Channel		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

<b>In Vitro</b>	DMSO : 10 mg/mL (23.56 mM; ultrasonic and adjust pH to 6 with HCl)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	<b>Preparing Stock Solutions</b>	1 mM	2.3562 mL	11.7808 mL	23.5616 mL
		5 mM	0.4712 mL	2.3562 mL	4.7123 mL
10 mM		0.2356 mL	1.1781 mL	2.3562 mL	
Please refer to the solubility information to select the appropriate solvent.					
<b>In Vivo</b>	<ol style="list-style-type: none"> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 40% PEG300 &gt;&gt; 5% Tween-80 &gt;&gt; 45% saline Solubility: ≥ 1 mg/mL (2.36 mM); Clear solution</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% (20% SBE-β-CD in saline) Solubility: ≥ 1 mg/mL (2.36 mM); Clear solution</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% corn oil Solubility: ≥ 1 mg/mL (2.36 mM); Clear solution</li> </ol>				

### BIOLOGICAL ACTIVITY

<b>Description</b>	MCHR1 antagonist 2 is an antagonist of melanin concentrating hormone receptor 1, with an IC <sub>50</sub> of 65 nM; MCHR1 antagonist 2 also inhibits hERG, with an IC <sub>50</sub> of 4.0 nM in IMR-32 cells <sup>[1]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	IC <sub>50</sub> : 65 nM (melanin concentrating hormone receptor 1), 4.0 nM (hERG in IMR-32 cells) <sup>[1]</sup>
<b>In Vitro</b>	MCHR1 antagonist 2 (Compound 30) is an antagonist of melanin concentrating hormone receptor 1, with an IC <sub>50</sub> of 65 nM. MCHR1 antagonist 2 has inhibitory effects on Ca <sup>2+</sup> flux, and hERG, with IC <sub>50</sub> s of 196 ± 30 nM and 4.0 ± 0.8 nM, respectively, in

---

IMR-32 cells<sup>[1]</sup>.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

---

## REFERENCES

---

[1]. Lynch JK, et al. Optimization of chromone-2-carboxamide melanin concentrating hormone receptor 1 antagonists: assessment of potency, efficacy, and cardiovascular safety. J Med Chem. 2006 Nov 2;49(22):6569-84.

---

**Caution: Product has not been fully validated for medical applications. For research use only.**

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: [tech@MedChemExpress.com](mailto:tech@MedChemExpress.com)

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA